WHAT IS CLAIMED IS:

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1. A method for recombinantly producing functional ataxia-telangiectasia (ATM) protein, comprising:

providing a viral vector comprising a gene encoding the ATM protein operably linked to a promoter;

infecting mammalian cells with said viral vector, wherein said mammalian cells produce functional ATM protein; and

isolating said functional ATM protein produced by said mammalian cells.

- 2. The method of Claim 1, wherein said viral vector comprising a gene encoding the ATM protein operably linked to a promoter is a vaccinia viral vector
- 3. The method of Claim 1, wherein said viral vector comprising a gene encoding the ATM protein operably linked to a promoter is a variola viral vector.
- 4. The method of Claim 1, wherein said viral vector comprising a gene encoding the ATM protein operably linked to a promoter is pSCAT
- 5. The method of Claim 1, wherein said promoter is a synthetic early/late viral promoter.
 - 6. The method of Claim 1, wherein said mammalian cells are human cells.
 - 7. The method of Claim 1, wherein said mammalian cells are HeLa cells.
- 8. The method of Claim' 1, wherein said mammalian cells are ATM-deficient cells.
 - 9. The method of Claim 1, wherein said mammalian cells are L3 cells.
- 10. The method of Claim 8, further wherein said ATM-deficient cells producing said functional ATM protein exhibit regain of ATM function.
- 11. The method of Claim 1 wherein isolating said functional ATM protein comprises binding an anti-ATM antibody to said ATM protein.
- 12. The method of Claim 1, where said gene encoding the ATM protein is modified to comprise a FLAG epitope.
- 13. The method of Claim 12, wherein isolating said functional ATM protein comprises binding an antibody specific for the FLAG epitope to said ATM protein.



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- 14. The method of Claim 1, wherein said functional ATM protein is produced at a level of greater than 2 ug substantially purified ATM protein per 300 grams fresh weight of host cells or host tissue.
- 15. The method of Claim 1, further wherein said functional ATM protein is capable of phosphorylating ATM substrates.
- 16. The method of Claim 15, wherein said substrates comprise p53 and PHAS-1.
- 17. A method for recombinantly producing a high yield of functional ataxia-telangiectasia (ATM) protein, comprising:

providing a vaccinia viral vector comprising a gene encoding the ATM protein operably linked to a promoter;

infecting mammalian cells with said vaccinia viral vector, wherein said mammalian cells produce functional ATM protein; and

isolating said functional ATM protein produced by said mammalian cells.

- 18. The method of Claim 17, wherein said high yield of functional ATM protein is greater than 2 ug substantially purified ATM protein per 300 grams fresh weight of mammalian cells.
 - 19. The method of Claim 17, wherein said mammalian cells are human cells.
- 20. The method of Claim 17, wherein said isolating said functional ATM protein comprises binding an anti-ATM antibody to the ATM protein.
- 21. The method of Claim 17, where said gene encoding the ATM protein is modified to comprise a FLAG epitope.
- 22. The method of Claim 21, wherein isolating said functional ATM protein comprises binding an antibody specific for the FLAG epitope to said ATM protein.

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